

What is claimed is:

1. A method for protecting cognitive function in a mammal comprising administering to the mammal a morphogen or nucleic acid encoding the morphogen.
2. A method for reducing memory dysfunction in a mammal, the method comprising the step of administering to the mammal a morphogen or a nucleic acid encoding the morphogen.
3. A method for treating dementia in a mammal, comprising the step of administering to the mammal a morphogen or a nucleic acid encoding the morphogen.
4. A method for treating a symptom associated with hippocampal tissue damage in a mammal, comprising the step of administering to the mammal a morphogen or a nucleic acid encoding the morphogen.
5. The method of claim 1, 2, 3 or 4 wherein said mammal is afflicted with or at risk of brain tissue damage associated with mechanical or chemical trauma, oxygen deprivation, glucose deprivation, a neurotoxin, a neurodegenerative disorder or dementia.
6. The method of claim 5 wherein said tissue damage results from ischemia.
7. The method of claim 1, 2, 3 or 4 wherein said mammal is a human.
8. The method of claim 6 wherein said human is at risk of or is afflicted with arterial occlusion cardiac arrest or stroke.
9. The method of claim 1, 2, 3 or 4 wherein said mammal is afflicted with or at risk of amnesia.
10. The method of claim 1, 2, 3 or 4 which said mammal is afflicted with or is at risk of Alzheimer's Disease, Pick Disease, Parkinson's Disease, amyotrophic lateral sclerosis,

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17. The method of claim 15 wherein said morphogen is any one of OP2, OP3, BMP2; BMP3; BMP4; BMP5; BMP6; BMP9; BMP-10, BMP-11, BMP-12, BMP-15, BMP-3b, DPP; Vg1; Vgr; 60A protein; GDF-1; GDF-3, GDF-5, GDF-6, GDF-7, GDF-8, GDF-9, GDF-10, GDF-11; and morphogenically active amino acid sequence variants thereof.
18. The method of claim 15 wherein said morphogen is noncovalently complexed with at least one morphogen pro domain.
19. A composition for protecting cognitive function and/or reducing cognitive dysfunction in a mammal, the composition comprising a morphogen in an amount sufficient to protect cognitive function and/or reduce cognitive dysfunction in said mammal.
20. The composition of claim 19 wherein said morphogen is dispersed in an aqueous solution.
21. The composition of claim 19 wherein said morphogen is disposed in a biodegradable, biocompatible matrix or binding agent.
22. The composition of claim 17 wherein said morphogen is disposed in a biocompatible microsphere.
23. A composition for protecting cognitive function and/or reducing cognitive dysfunction in a mammal, the composition comprising cultured cells competent to express a morphogen in an amount sufficient to protect cognitive function and/or reduce cognitive dysfunction in said mammal.
24. The composition of claim 23 wherein said cells are disposed in a porous, biocompatible material.
25. A composition for protecting cognitive function and/or reducing cognitive dysfunction in a mammal, the composition comprising a recombinant nucleic acid comprising a DNA sequence encoding a morphogen and a promoter in operative association therewith, in an

26. A kit for protecting cognitive function and/or reducing cognitive dysfunction in a mammal, the kit comprising

1. The first part of the paper is devoted to the study of the asymptotic behavior of the solutions of the system (1) as $\epsilon \rightarrow 0$. It is shown that the solutions of the system (1) converge to the solutions of the system (2) in the sense of the L^2 -norm.

add E_4 add B_1 \rightarrow add E_4